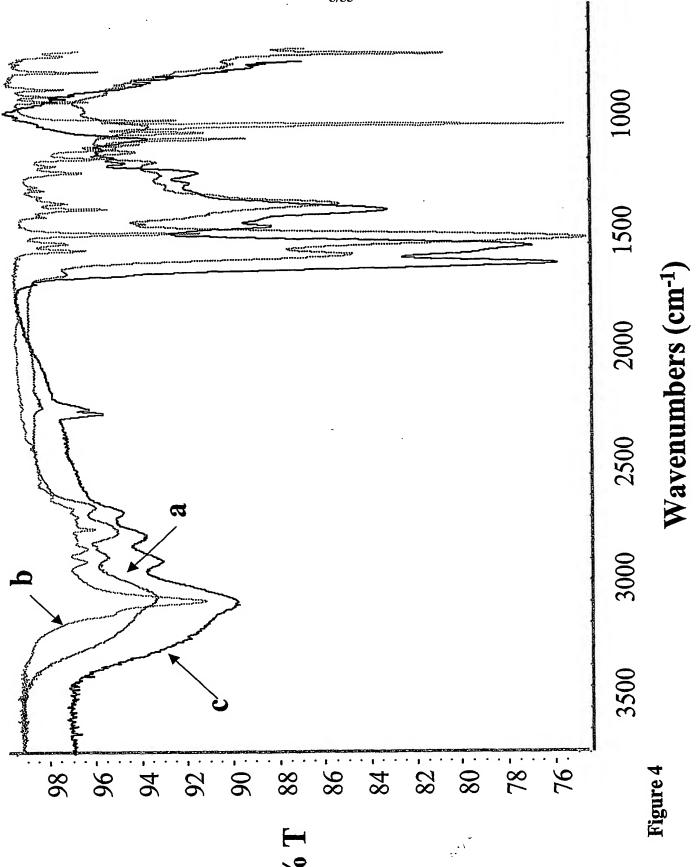
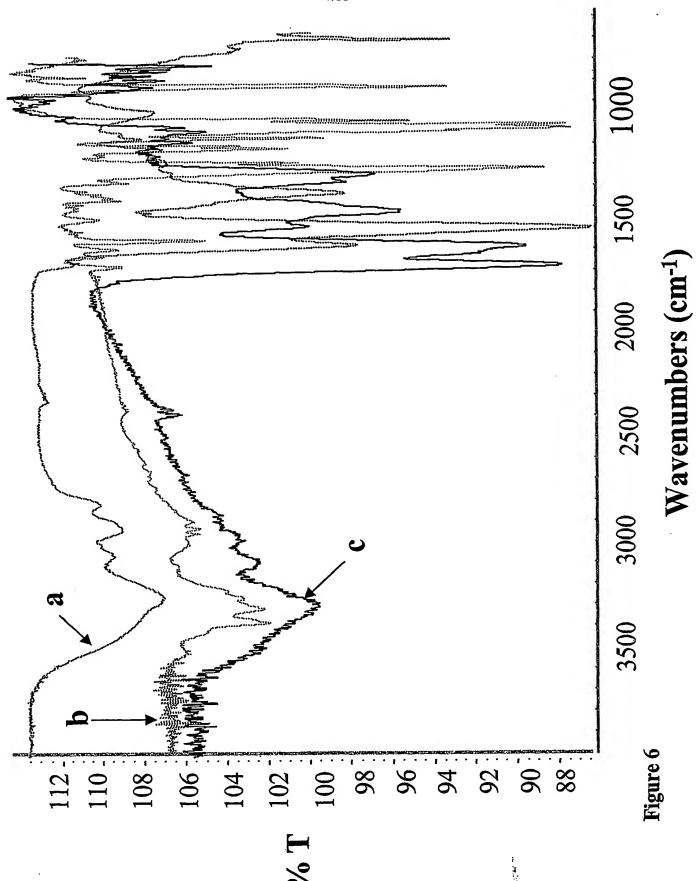


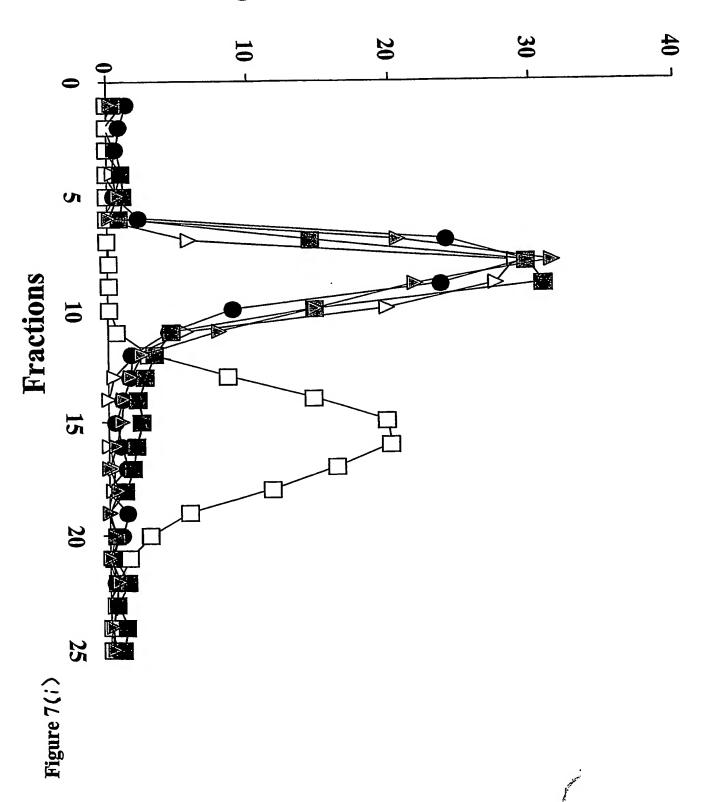
percentage (%) of 14C-glucosamine

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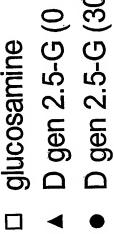




Percentage (%) of ¹4C-glucosamine







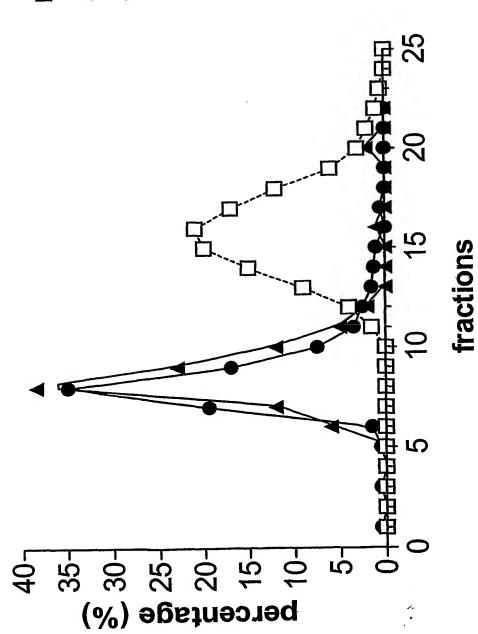
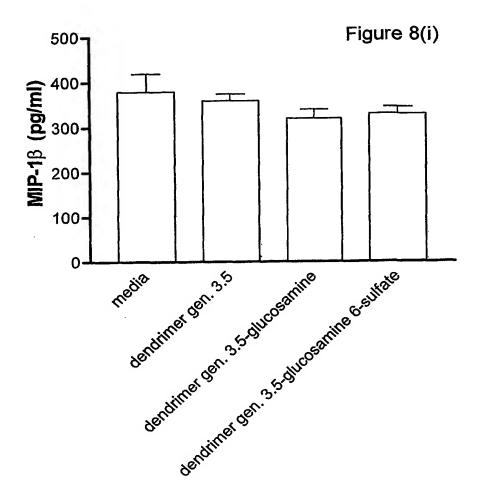
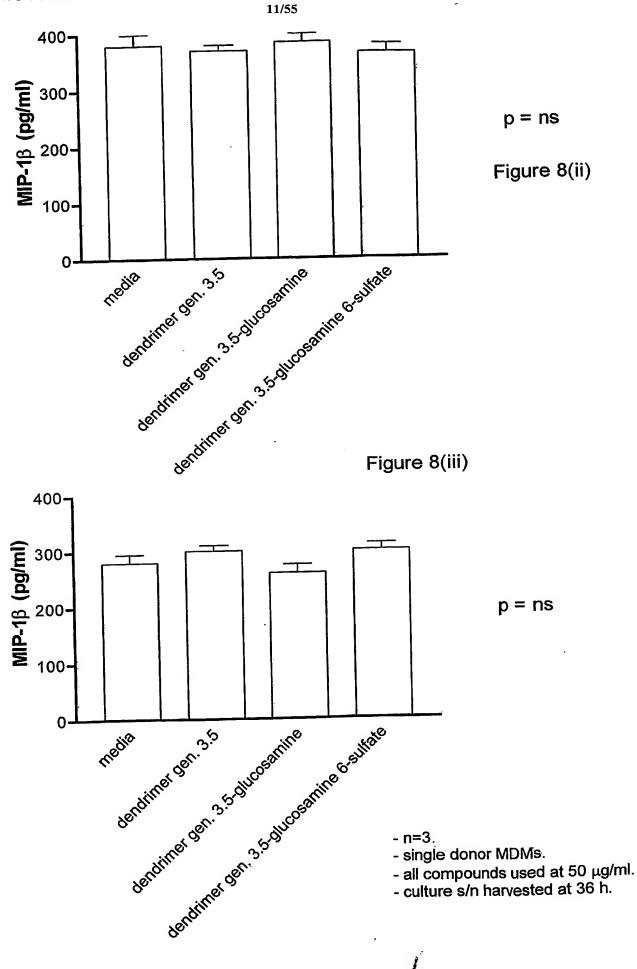


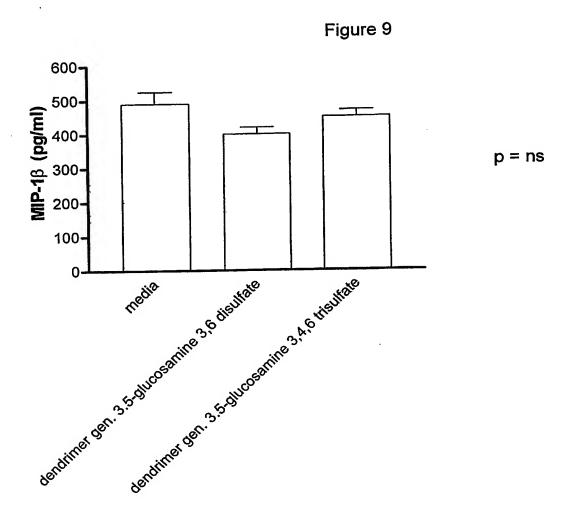
Figure 7(ii): PD-10 column of dendrimers gen 2.5-glucosamine after synthesis [D gen 2.5-G (0 d)] and after 30 days of storage [D gen 2.5-G (30 d)]



- n = 3. Results shown as mean \pm sem. p = ns.
- single donor PBMN cells.
- all compounds used at 50 μg/ml.
- culture supernatants harvested for MIP-1 $\!\beta$ at 36 h.

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- n = 3.
- single donor MDMs.
- all compounds used at 50 μg/ml.
- culture supernatants harvested for MIP-1 β after 36 h.

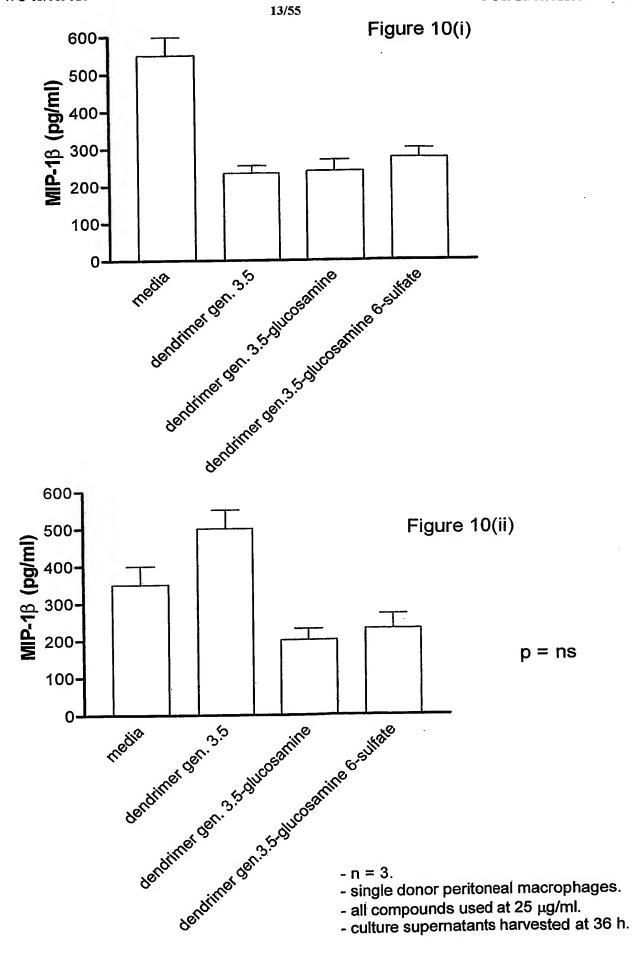
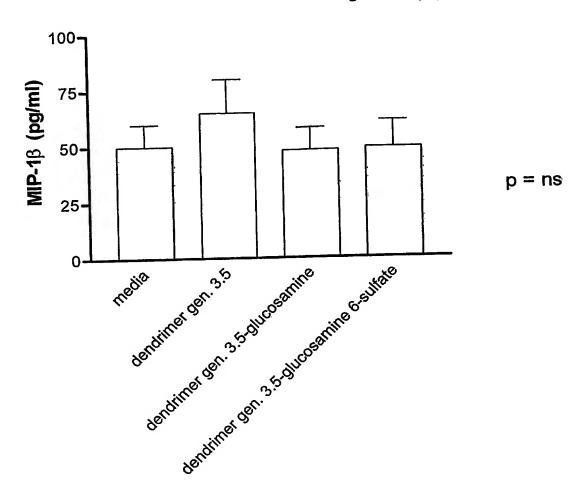
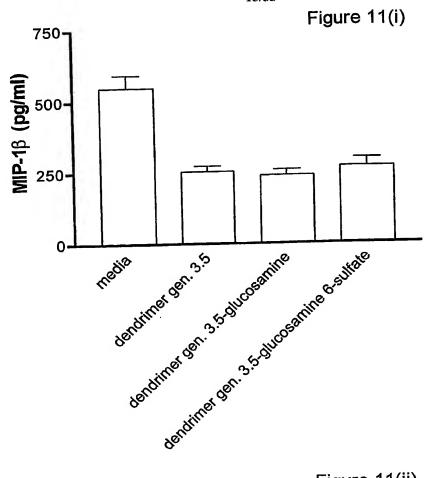
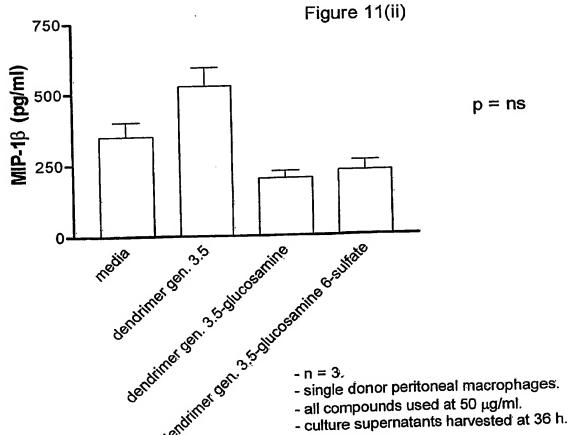


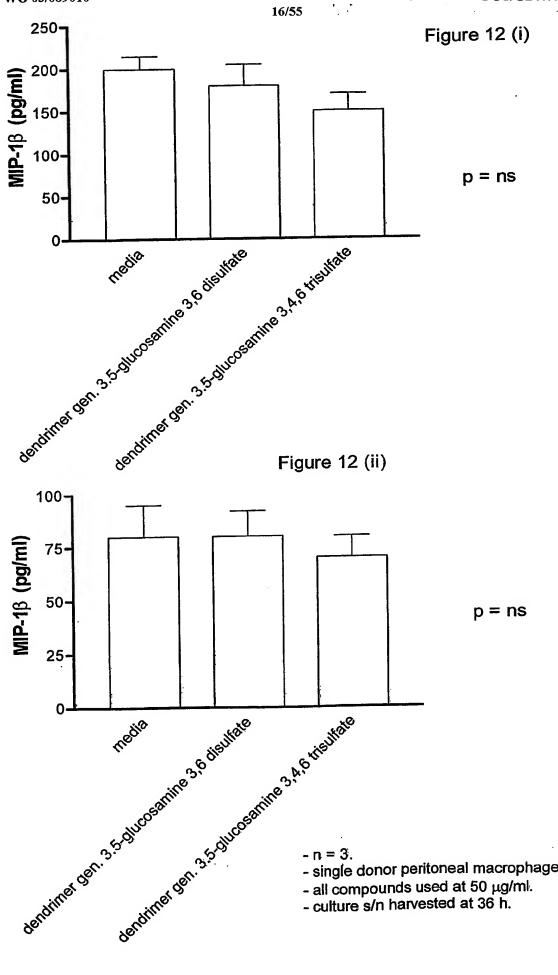
Figure 10(iii)



- -n = 3.
- single donor peritoneal macrophages.
- all compounds used at 25 $\mu\text{g/ml}.$
- culture supernatants harvested for MIP-1 β at 72 h.

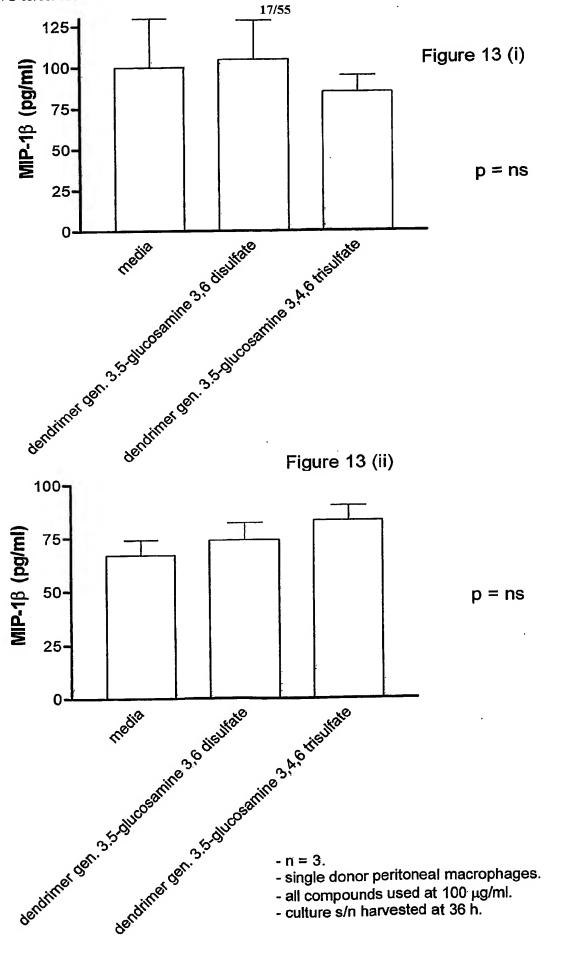


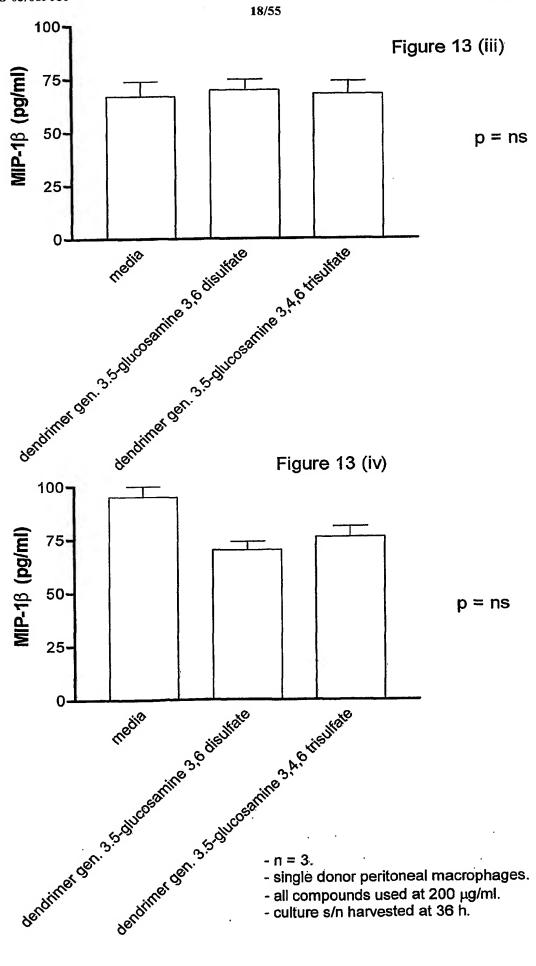


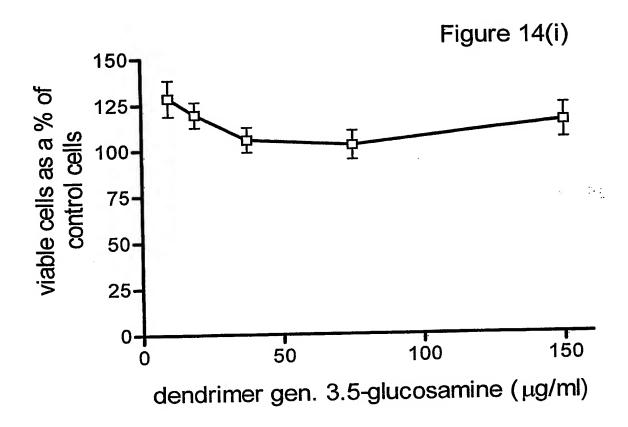


- single donor peritoneal macrophages.

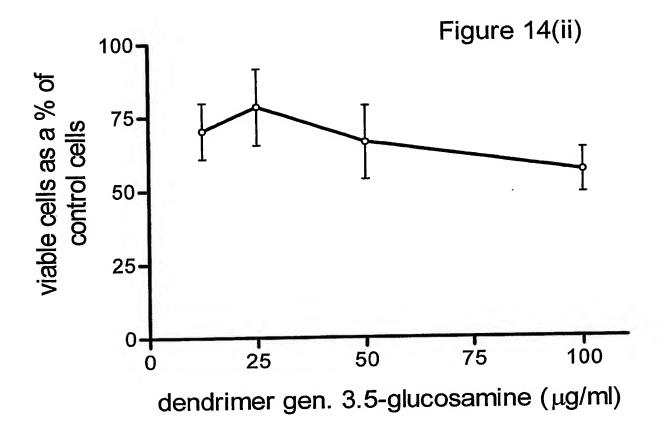
- all compounds used at 50 µg/ml. - culture s/n harvested at 36 h.



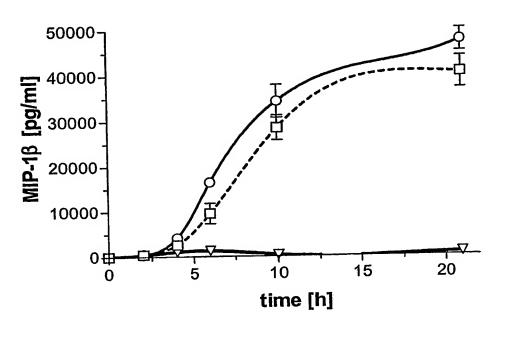




- -n = 3.
- single donor MDMs.
- MTT assay.

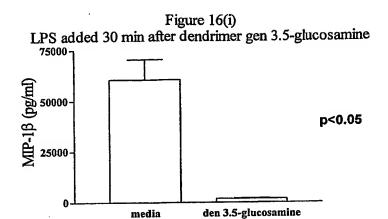


- n = 3. HUVECs.
- MTT assay.



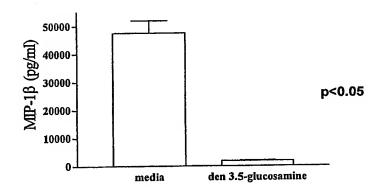
- □ LPS (5 ng/ml)
- o LPS (10 ng/ml)

Figure 15: Time course of the release of MIP-1 β into cell free culture supernatants after the addition of LPS at a final concentration of 5 ng/ml & 10 ng/ml.



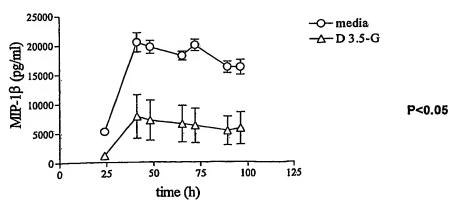
- n = 4 - mean ± sem - D 3.5-G at 100 μg/ml - LPS at 5 ng/ml - harvest at 21 h

Figure 16(ii)
LPS added 1 h after
dendrimer gen 3.5-glucosamine



- n = 4
- mean ± sem
- D 3.5-G at 100 μg/ml
- LPS at 5 ng/ml
- harvest at 21 h

Figure 16(iii)
LPS was added 24 h after dendrimer gen 3.5-glucosamine





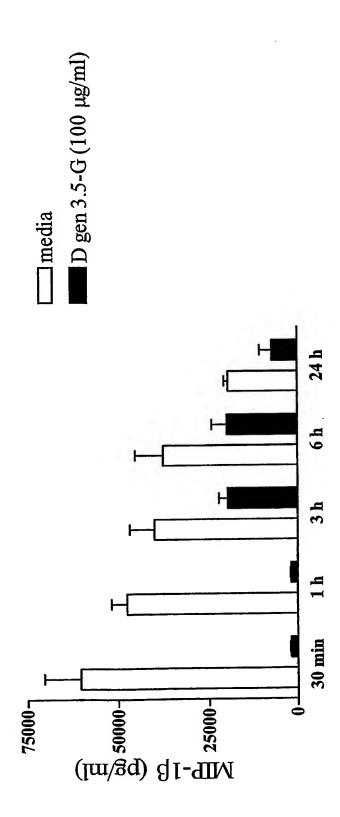
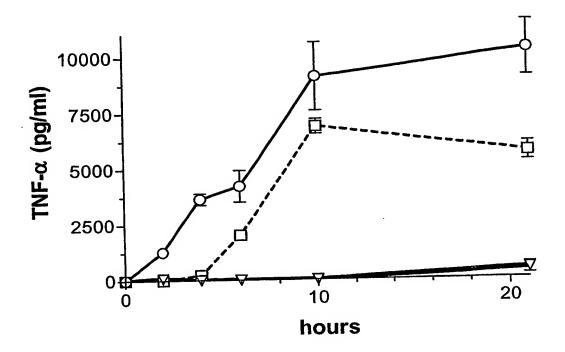


Figure 16(iv)

- Release of MIP-1 β from single donor PBMN cells. n = 4.
- LPS (5 ng/ml) was added at various time points ranging from
- Culture supernatants harvested 21 h after the addition of LPS for MIP-1 β . 30 min to 24 h after dendrimer gen. 3.5-glucosamine (100 µg/ml).
- (D gen 3.5-G: dendrimer gen 3.5-glucosamine)

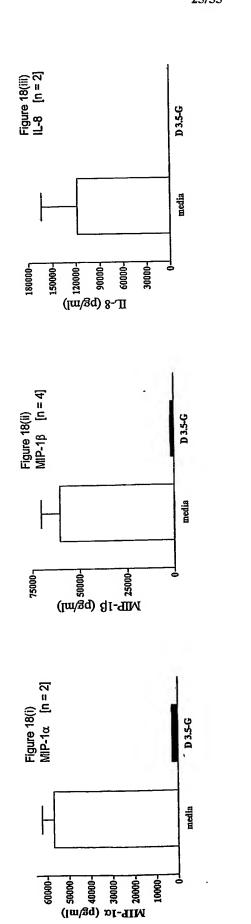
PCT/GB03/01133

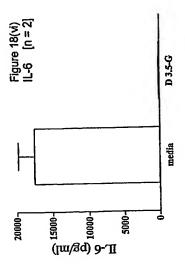


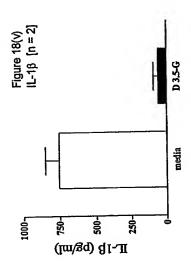
- ▽ control
- □ LPS (5 ng/ml)
- o LPS (10 ng/ml)

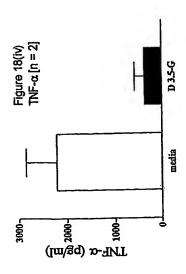
Figure 17: Time course of the release of TNF- α into cell free culture supernatants following the addition of LPS at a final concentration of 5 ng/ml and at 10 ng/ml.

Figure 18: LPS (5 ng/ml) was added 30 min after dendrimer gen 3.5-glucosamine (100 μg/ml) and culture supernatants harvested at 21 h. (P<0.05 for all 6 figures)









D35-G

media.

Figure 19: LPS (5 ng/ml) was added 30 min after dendrimer gen 3.5-glucosamine (200 μg/ml) and culture supernatants harvested at 25 h. (P<0.05 for all groups)

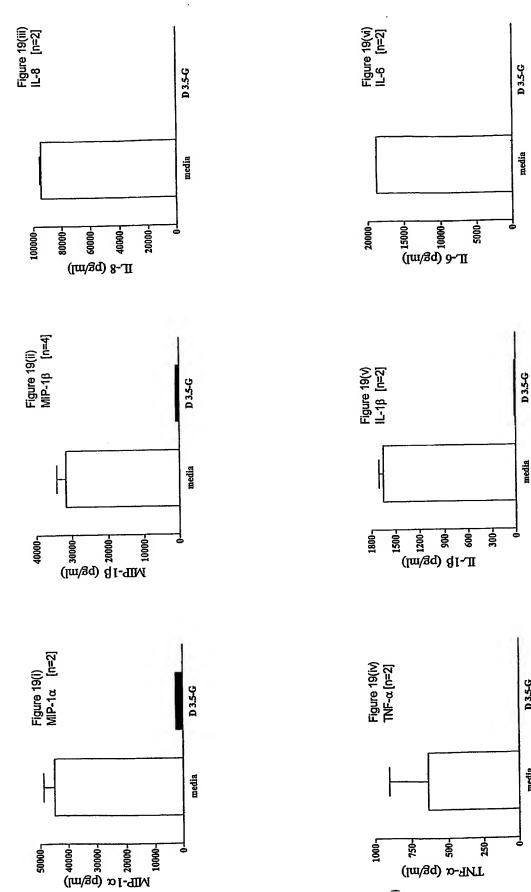
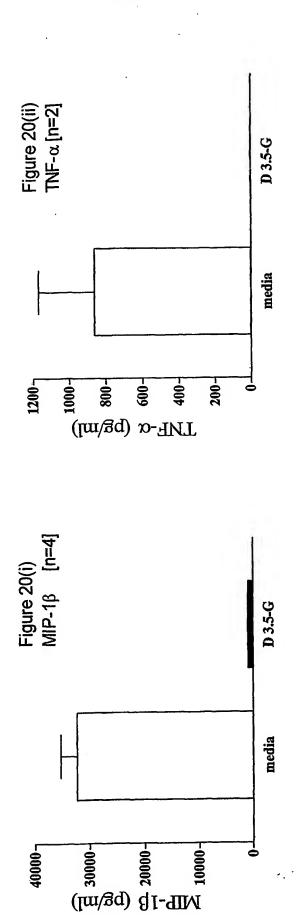
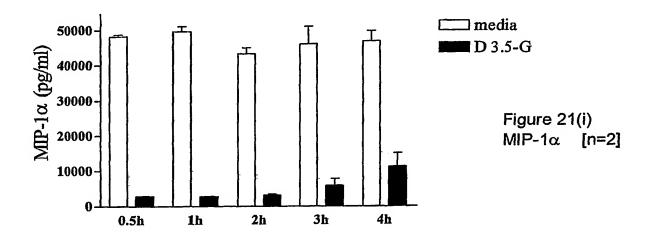


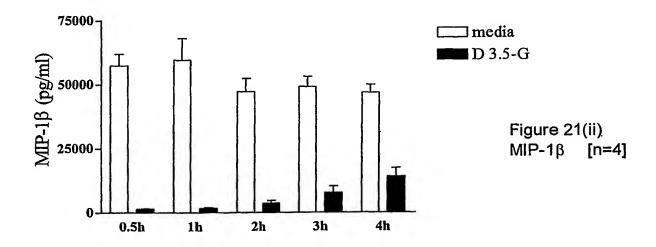
Figure 20: LPS (5 ng/ml) added 1 hour after dendrimer gen 3.5-glucosamine (200 μg/ml) and culture supernatants harvested at 25 h. (P<0.05 for all groups)



mean ± sem

Figure 21(i): Dendrimer gen 3.5-glucosamine ((D 3.5 G at 100 μ g/ml) added 30 min or 1 h or 2 h or 3 h or 4 h after LPS (5 ng/ml). Culture supernatants harvested at 21 h.





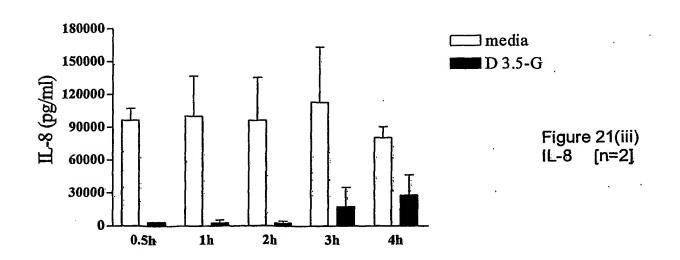
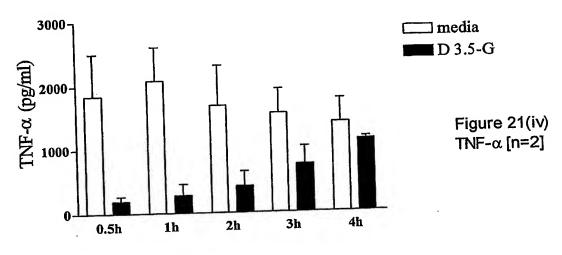
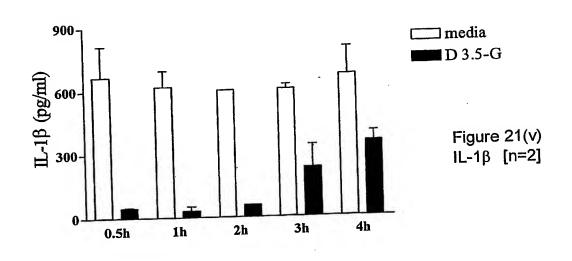
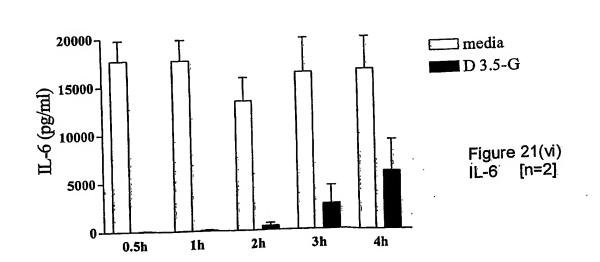


Figure 21(ii): Dendrimer gen 3.5-glucosamine (100 μ g/ml) added 30 min or 1 h or 2 h or 3 h or 4 h after LPS (5 ng/ml). Culture supernatants harvested at 21 h.







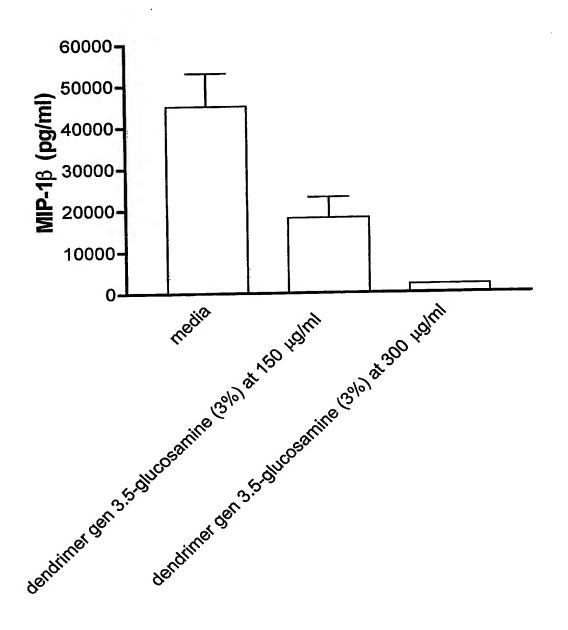


Figure 22(i): Release of MIP-1 β from single donor PBMN cells. LPS (5ng/ml) was added 30 min after the dendrimer generation 3.5-glucosamine (3% loading) at a concentration of either 150 μ g/ml or 300 μ g/ml. Culture supernatants were harvested at 22 h (n=1).

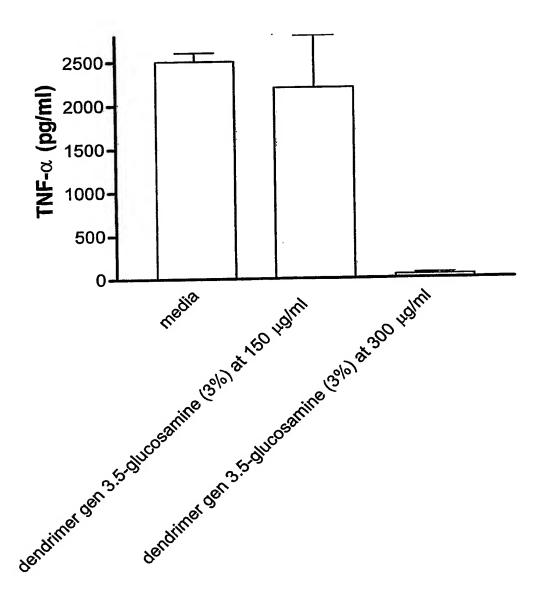
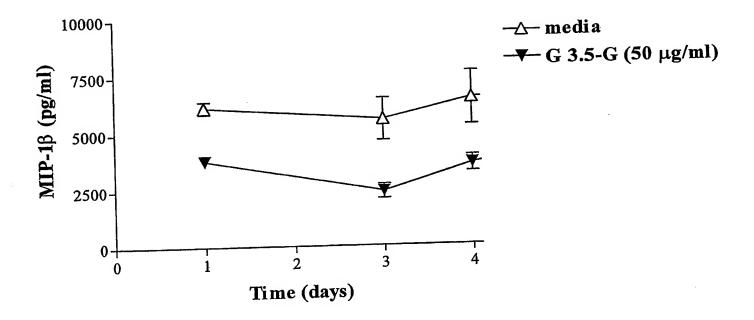


Figure 22(ii): Release of TNF- α from single donor PBMN cells. LPS (5ng/ml) was added 30 min after the dendrimer generation 3.5-glucosamine (3% loading) at a concentration of either 150 μ g/ml or 300 μ g/ml. Culture supernatants were harvested at 22 h (n=1).

Figure 23(i)
PBMN cells from 3 donors mixed [n=2]



33/55

Figure 23(ii)
PBMN cells from 3 donors mixed [n=2]

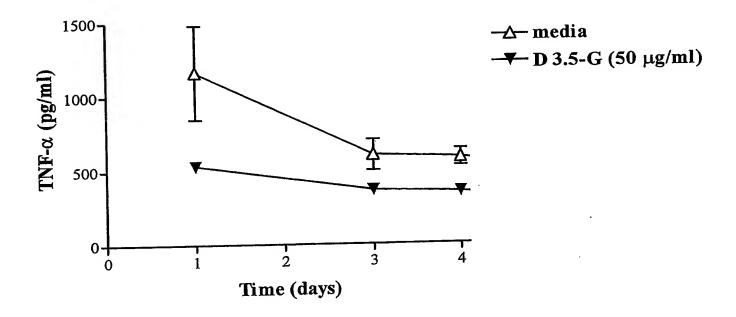


Figure 23(iii)
PBMN cells from 2 donors mixed [n=2]

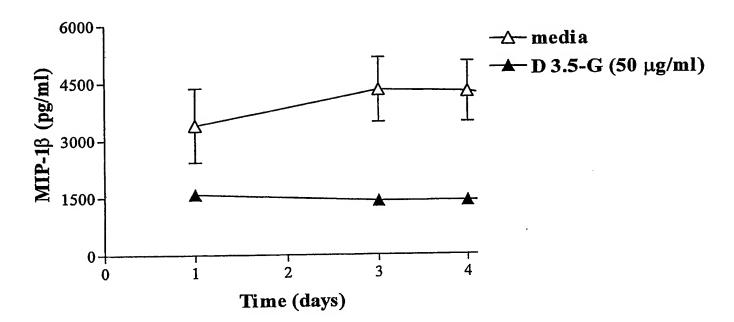
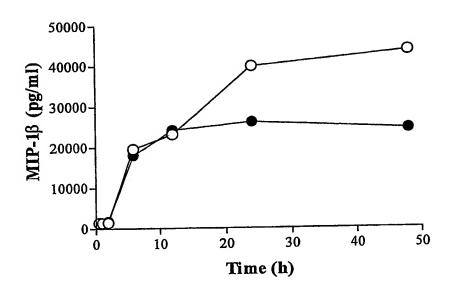


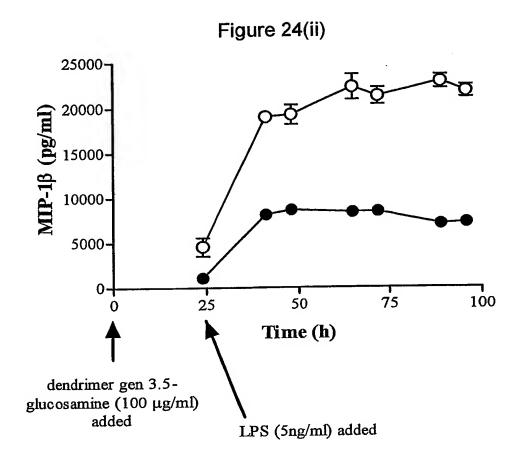
Figure 24(i) PBMN cells from 2 donors mixed [n=2]



-O-LPS + media

--- LPS + dendrimer gen 3.5-glucosamine at 100 μg/ml

Donor PBMN cells from 2 individuals mixed, Dendrimer gen 3.5-glucosamine (100 µg/ml) followed immediately by LPS at 10 ng/ml).



-O-LPS (5 ng/ml) + media

-- LPS (5 ng/ml) + dendrimer gen 3.5-glucosamine at 100 μg/ml

PBMN cells from 2 individuals were mixed and the dendrimer gen 3.5-glucosamine added at 100 μ g/ml. 24 hours later, LPS was added at 5 ng/ml (n=2).

Figure 25(i) PBMN cell viability when cultured with compounds for up to 5 days.

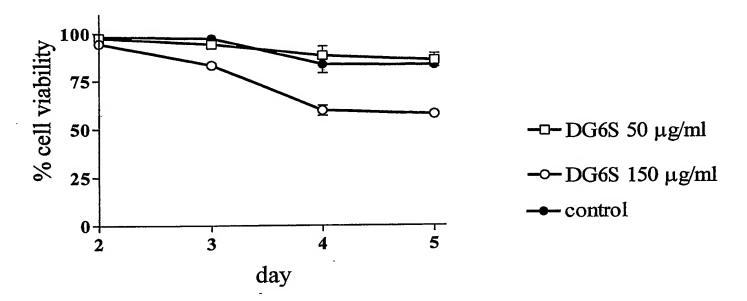


Figure 25(ii)
PBMN cell counts when cultured with compounds for up to 5 days.

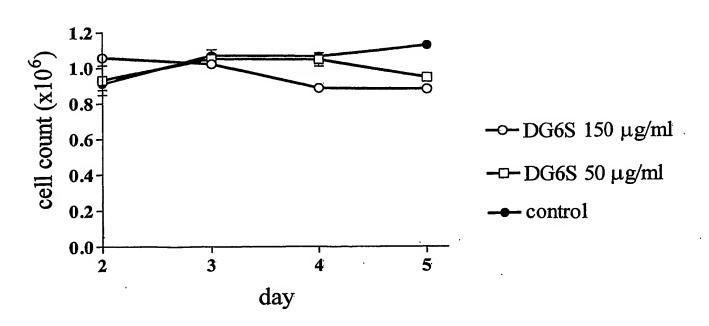
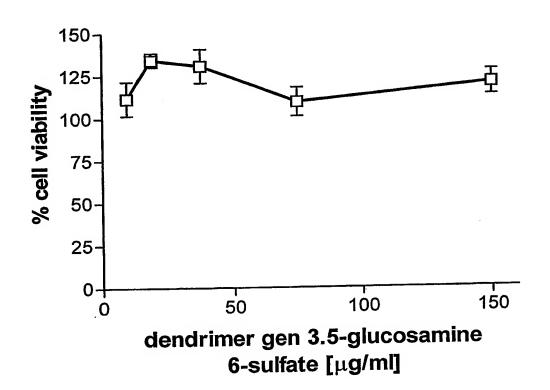
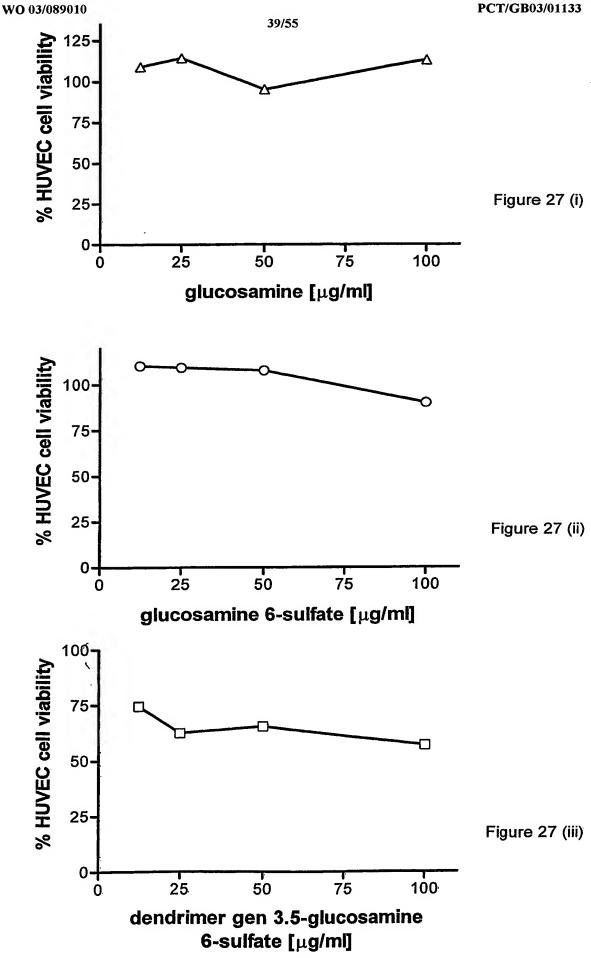


Figure 26: Cell viability of MDMs when cultured with dendrimer gen 3.5-glucosamine 6-sulfate over a period of 5 days





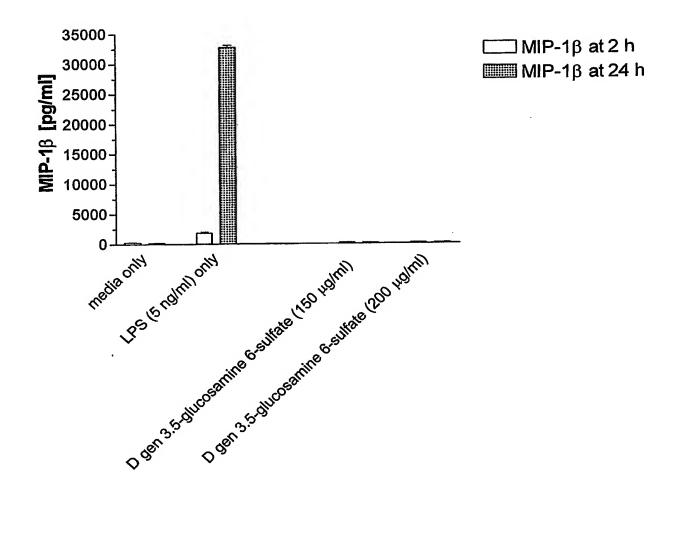


Figure 28(i): MIP-1 β release from single donor PBMN cells when treated with dendrimer gen 3.5-glucosamine 6-sulfate at 150 μ g/ml or 200 μ g/ml for 30 min before the addition of LPS at 5 ng/ml.

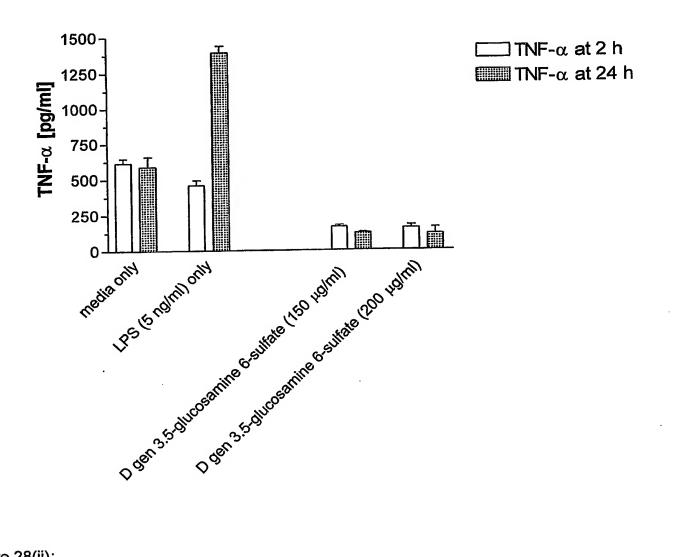


Figure 28(ii): TNF- α release from single donor PBMN cells when treated with dendrimer gen 3.5–glucosamine 6-sulfate at 150 μ g/ml or 200 μ g/ml for 30 min before the addition of LPS at 5 ng/ml.

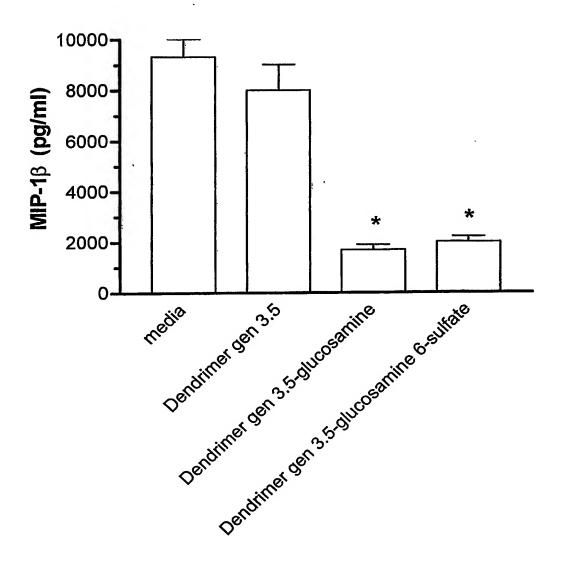
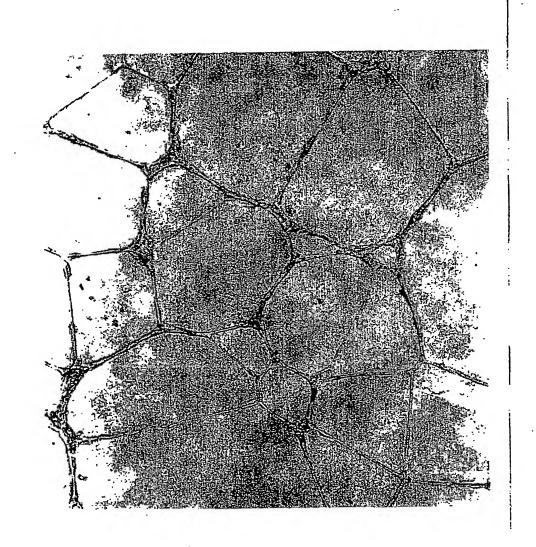


Figure 29: MDMs from four donors were pooled for 24 h and each of the compounds shown then added at a concentration of 125 μ g/ml. Cell free culture supernatants were then harvested at 36 h for measurement of MIP-1 β . A reduction in MIP-1 β was seen when dendrimer gen 3.5 glucosamine or dendrimer gen 3.5-glucosamine 6-sulfate was present.

^{*} p<0.05 compared to the media control

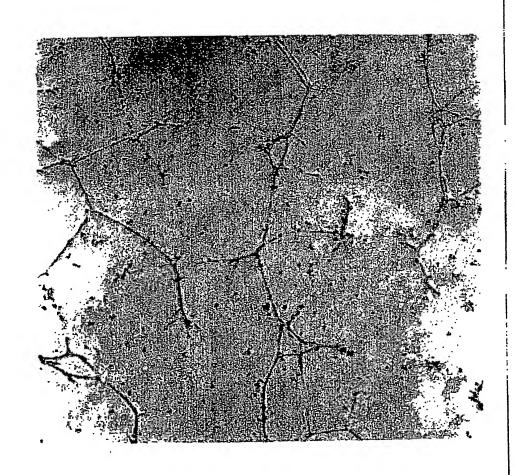
A visual analogue scale was used to determine the extent of tube formation and scored on a scale from 0 (all cells remain single) to 4 (all cells involved in Figure 30: Endothelial microtubule formation by HUVECs on Matrigel (x 40 magnification). tubular structures) as described in the text.

Control well



A visual analogue scale was used to determine the extent of tube formation and scored on a scale from 0 (all cells remain single) to 4 (all cells involved in Figure 31: Endothelial microtubule formation by HUVECs on Matrigel (x 40 magnification). tubular structures) as described in the text.

Dendrimer gen. 3.5-glucosamine 6-sulfate at 12.5 µg/ml



A visual analogue scale was used to determine the extent of tube formation and scored on a scale from 0 (all cells remain single) to 4 (all cells involved in Figures 32: Endothelial microtubule formation by HUVECs on Matrigel (x 40 magnification). tubular structures) as described in the text.

Dendrimer gen 3.5-glucosamine 6-sulfate at 50 µg/ml

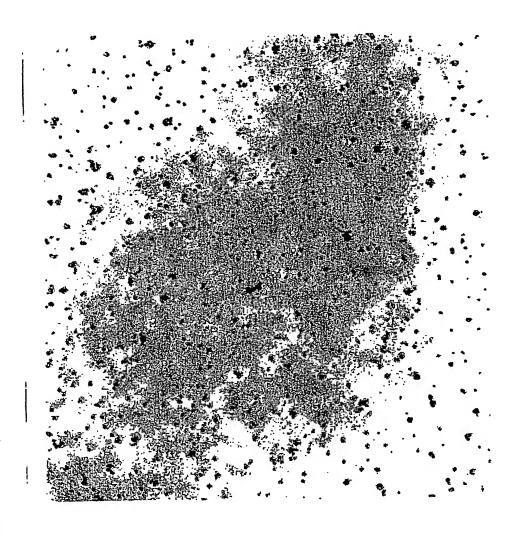
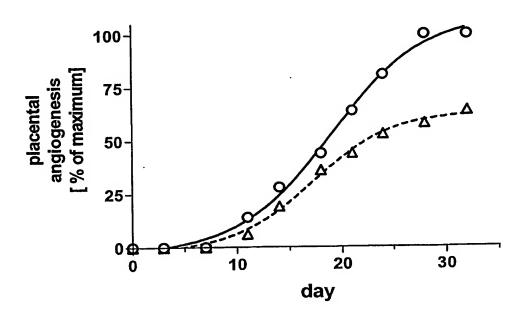


Figure 33: Dendrimer gen. 3.5 glucosamine 6-sulfate reduced the rate of new blood vessel formation in an *in vitro* human placental angiogenesis assay.



- o Control
- Δ Dendrimer gen. 3.5 glucosamine 6-sulfate (50 μ g/ml)

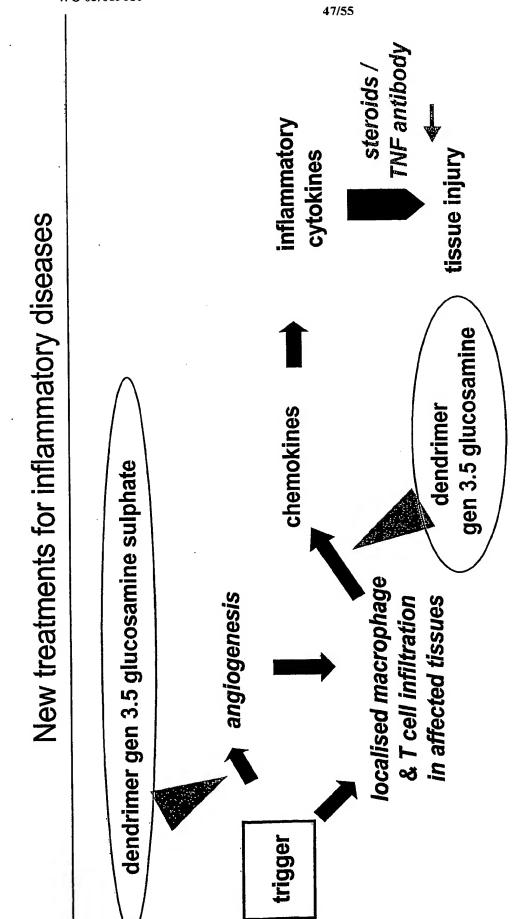
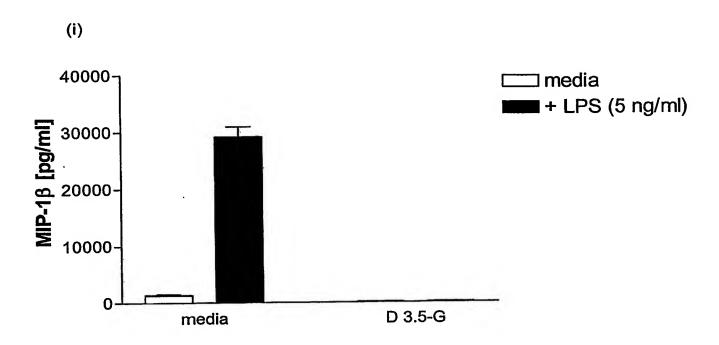


Figure 34

Figure 35: Release of MIP-1 β and TNF- α from single donor monocyte derived dendritic cells that were exposed to LPS (5 ng/ml) in the absence and the presence of dendrimer gen. 3.5 glucosamine (D 3.5-G) at 200 μ g/ml.



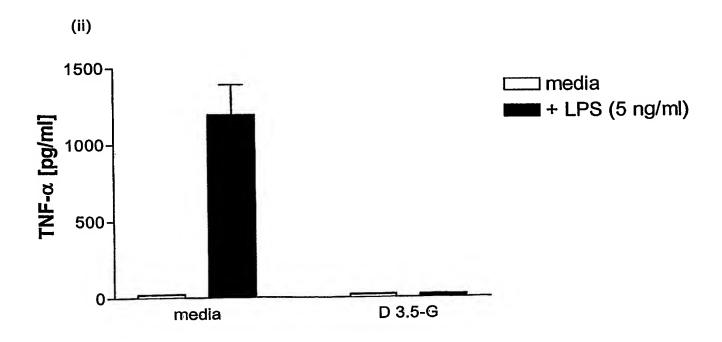
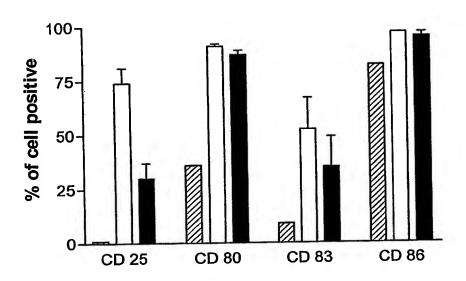


Figure 36: Effect of dendrimer gen. 3.5-glucosamine (200 μ g/ml) on the upregulation of CD 25, CD 80, CD 83 and CD 86 on monocyte derived dendritic cells by LPS (5 ng/ml).

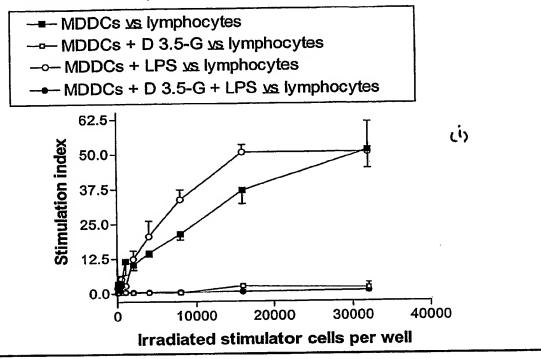


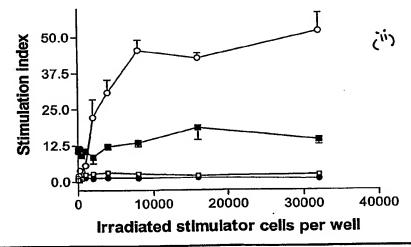
Media control

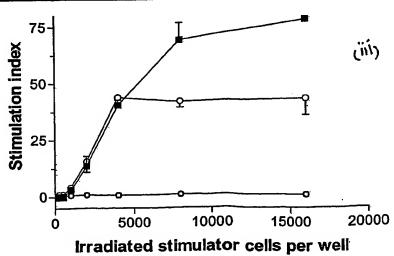
____LPS (5 ng/ml)

Dendrimer gen. 3.5-glucosamine (200 μg/ml) + LPS (5 ng/ml)

Figure 37: Effect of dendrimer gen. 3.5 glucosamine (D 3.5-G at 200 $\mu g/ml$) on the allogenic mixed lymphocyte reaction using monocyte derived dendritic cells (MDDCs) in the absence & the presence of LPS (5 ng/ml).

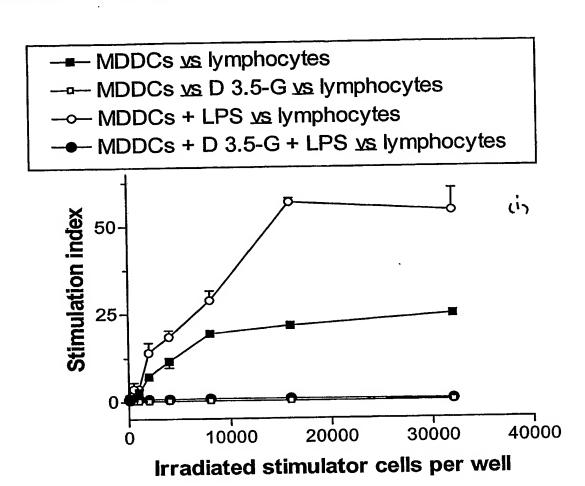






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Figure 38: Effect of dendrimer gen. 3.5 glucosamine (D 3.5-G at 200 μ g/ml) on the allogeneic mixed lymphocyte reaction using monocyte derived dendritic cells (MDDCs) in the absence and the presence of LPS (20 ng/ml).



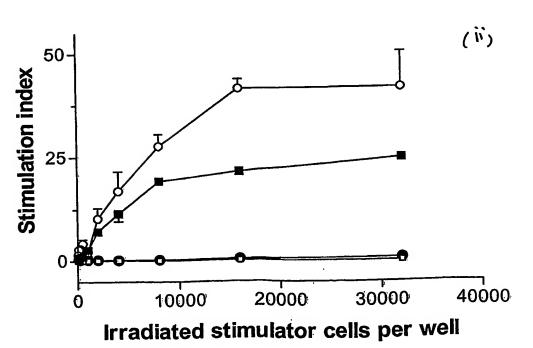
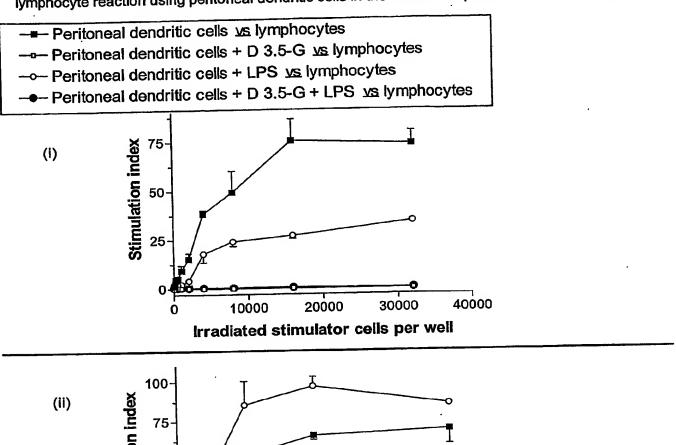
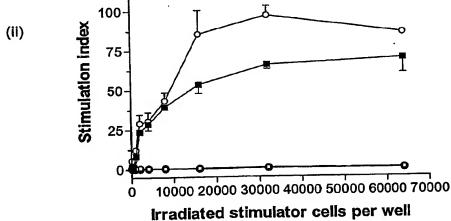


Figure 39: Effect of dendrimer gen. 3.5 glucosamine (D 3.5-G at 200 μ g/ml) on the allogeneic mixed lymphocyte reaction using peritoneal dendritic cells in the absence & presence of LPS (5 ng/ml).





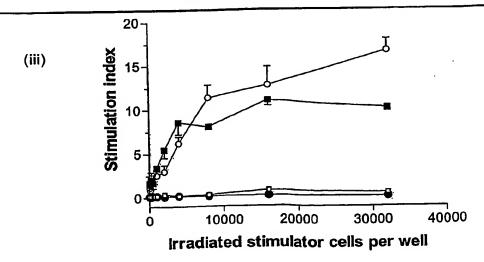
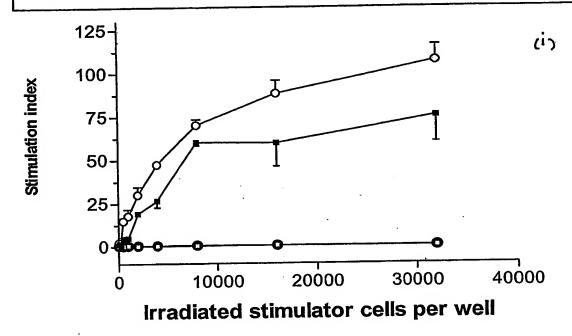


Figure 40: Effect of dendrimer gen. 3.5 glucosamine (D 3.5-G at 200 μ g/ml) on the allogeneic mixed lymphocyte reaction using peritoneal dendritic cells in the absence and the presence of LPS (20 ng/ml).

- Peritoneal dendritic cells vs lymphocytes
- —□— Peritoneal dendritic cells + D 3.5-G <u>vs</u> lymphocytes
- Peritoneal dendritic cells + D 3.5-G + LPS <u>vs</u> lymphocytes



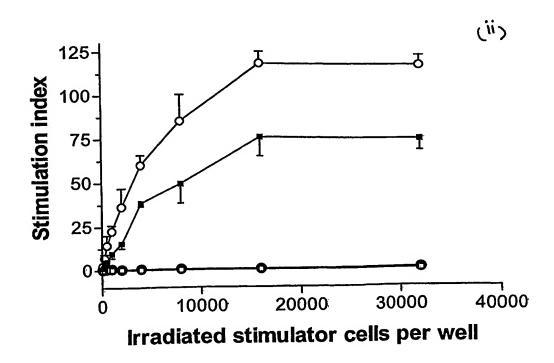
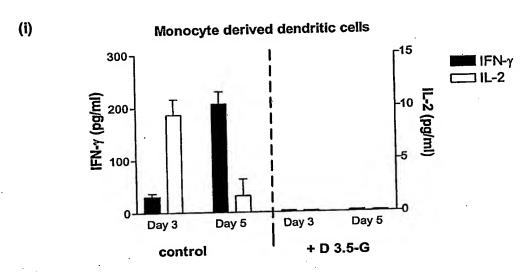
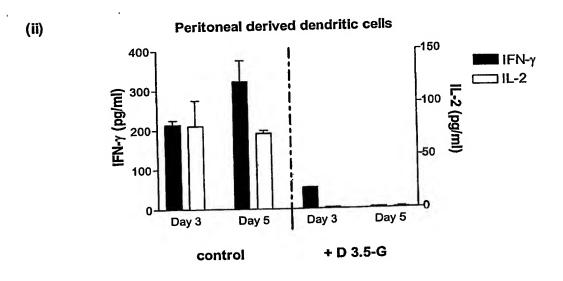


Figure 41:

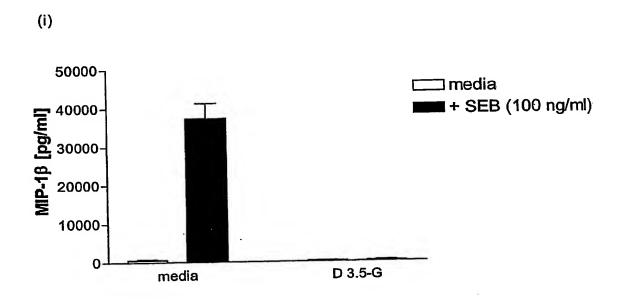


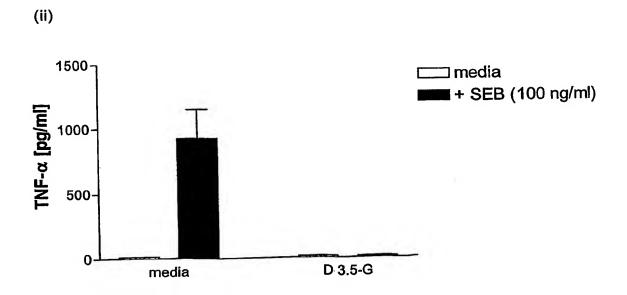
32,000 monocyte derived dendritic cells were stimulated with LPS (20 ng/ml) for 21 h, washed and incubated with 100,000 allogeneic lymphocytes. Cell free supernatants were then analysed at days 3 and 5. (D 3.5-G = dendrimer gen. 3.5 glucosamine)



32,000 peritoneal dendritic cells were stimulated with LPS (20 ng/ml) for 21 h, washed and incubated with 100,000 allogeneic lymphocytes. Cell free culture supernatants were then analysed at days 3 and 5.

Figure 42: Release of MIP-1 β and TNF- α from single donor PBMN cells that were exposed to SEB (100 ng/ml) in the absence and the presence of dendrimer gen 3.5 glucosamine (D 3.5-G) at 200 μ g/ml.





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